

SUB
DP
COO+
C3
COO1

obtaining an mRNA;
reverse transcribing the mRNA into cDNA with reverse transcriptase without RNase H activity
so that a cDNA-mRNA complex is formed;
degrading the mRNA from the cDNA-mRNA complex to form a linear cDNA;
ligating the ends of said linear cDNA to form a circular cDNA;
introducing first and second sequence specific primers to said circular cDNA; and
initiating a primer extension amplification reaction to increase copy number of said circular
cDNA.

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C4 D1

6. (Amended)

The method of claim 1 further comprising the step of:
harvesting said amplified cDNA.

9. (Amended)

C5

The method of claim 1 wherein said first and second primers are designed to hybridize to
from about 4 to about 35 contiguous bases from a sequence known or suspected to be present in
said circular cDNA.

10. (Amended)

The method of claim 1 wherein said first primer comprises a 3' end of the same which is
toward the 5' end of the circular cDNA.

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11. (Amended)

The method of claim 1 wherein one of said primers comprises a 3' end of the same which
is toward the 3' end of said circular cDNA.

12. (Twice Amended)

A method for amplifying a cDNA, including the 5' and 3' ends, comprising:
obtaining an mRNA;
contacting the mRNA with reverse transcriptase without RNase H so that a cDNA-mRNA
complex is formed, and degrading the mRNA from the cDNA-mRNA complex to form a
linear cDNA;
circularizing said linear cDNA;
contacting the circularized cDNA with first and second sequence specific primers;
and

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004,
introducing a polymerase and a supply of nucleotide bases to said circularized cDNA so that an amplification reaction occurs, wherein said region of said cDNA outside of said first and second primers including the 3' and 5' ends of said cDNA is amplified.

15. (Amended)

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The method of claim 1 wherein said forward and reverse primers are designed to hybridize to from about 4 to about 35 contiguous bases from a sequence known or suspected to be present in said circular cDNA.

16. (Amended)

The method of claim 1 wherein said one of said primers comprises a 3' end of the same which is toward the 5' end of the circular cDNA.

17. (Amended)

The method of claim 1 wherein one of said primers comprises a 3' end of the same which is toward the 3' end of said circular cDNA.

26. (Amended)

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A method for amplifying a cDNA comprising:
obtaining an mRNA;
reverse transcribing the mRNA into cDNA with reverse transcriptase without RNase H activity so that a cDNA-mRNA complex is formed;
degrading the mRNA from the cDNA-mRNA complex to form a linear cDNA;
ligating the ends of said linear cDNA to form a circular cDNA;
introducing first and second sequence specific primers to said circular cDNA, wherein said primers are degenerate primers; and
initiating a primer extension amplification reaction to increase copy number of said circular cDNA.

27. (Amended)

A method for amplifying a cDNA comprising:
obtaining an mRNA;
reverse transcribing the mRNA into cDNA with reverse transcriptase without RNase H activity so that a cDNA-mRNA complex is formed;
degrading the mRNA from the cDNA-mRNA complex to form a linear cDNA;
ligating the ends of said circular cDNA to form a circular cDNA;